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FILE 'REGISTRY' ENTERED AT 10:10:49 ON 13 MAY 2008
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     FILE 'LREGISTRY' ENTERED AT 08:32:12 ON 13 MAY 2008
               ACT LOE060/0
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L1
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     FILE 'HCAPLUS' ENTERED AT 09:33:38 ON 13 MAY 2008
L4
          11867 S LEHMANN ?/AU
L5
             65 S LONSKY ?/AU
L6
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                SEL L6 1 RN
    FILE 'REGISTRY' ENTERED AT 09:36:03 ON 13 MAY 2008
L7
             51 S E1-E51
             34 S L7 AND SI/ELS
1.8
L9
            11 S L8 AND RSD/FA
             4 S L9 AND 3/NRRS
L10
             7 S L9 NOT L10
L11
    FILE 'HCAPLUS' ENTERED AT 09:52:32 ON 13 MAY 2008
L12
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    FILE 'LREGISTRY' ENTERED AT 10:01:17 ON 13 MAY 2008
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STR L13

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L17 51 S L15 FUL

DEL LOE060/Q
SAV L17 LOE060/A

FILE 'HCAPLUS' ENTERED AT 10:06:56 ON 13 MAY 2008

L18 25 S L17 L19 9 S L18 AND L12 L20 9 S L19 OR L12

L21 16 S L18 NOT L20

L22 8 S 1840-2003/PY,PRY,AY AND L20 L23 12 S 1840-2003/PY,PRY,AY AND L21

FILE 'REGISTRY' ENTERED AT 10:10:49 ON 13 MAY 2008

=> D L17 QUE STAT L15 STR

VAR G1=22/23/24 REP G2=(1-20) C NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 24

STEREO ATTRIBUTES: NONE

L17 51 SEA FILE=REGISTRY SSS FUL L15

100.0% PROCESSED 918 ITERATIONS SEARCH TIME: 00.00.01 => FILE HCAP

FILE 'HCAPLUS' ENTERED AT 10:11:36 ON 13 MAY 2008 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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=> D L22 1-8 BIB ABS HITSTR HITIND

- L22 ANSWER 1 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN
- 2005:472204 HCAPLUS AN

143:35205 DN

- Antireflective coatings for via fill and photolithography TT applications and
- methods of preparation thereof
- IN Li, Bo; Kennedy, Joseph; Iwamoto, Nancy; Lu, Victor; Leung, Roger; Fradkin, Mark A.; Hussein, Makarem A.; Goodner, Michael D.
- PA Honeywell International Inc., USA
- SO PCT Int. Appl., 120 pp. CODEN: PIXXD2

DT Patent

LA English

FAN CNT 1

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|------------|------------|------|------|------|------|------|----------|-------|-------|-----------------|-----|------|--------|-----|-------|-------|---|
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| ΡI | WO 2005 | 0496 | 81 | | A2 | | 20050602 | | | WO 2004-US38517 | | | | | | | |
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| | WO 2005 | 0496 | 81 | | A3 | | 2006 | 0420 | | | | | | | | | |
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| GD, | | | | | | | | | | | | | | | | | |
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             SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
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             NE, SN, TD, TG
     US 20050171277
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                                20050804
                                           US 2003-717028
20031118 <--
     EP 1695142
                                20060830 EP 2004-811280
                          A 2.
20041117 <--
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     CN 1902546
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                                            JP 2006-541340
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PRAI US 2003-717028
                          Α
                                20031118 <--
     WO 2004-US38517
                                20041117
                          W
     An antireflection coating that contains an optical absorber which
AB
     absorbs strongly and uniformly in the UV regions, keeps the
     photoresist from falling over and expanding outside or inside the
     resist region, is imperious to developers and methods of production
     of SOG antireflective coatings, satisfies goals of etching or
     stripping selectivity, minimizes filling bias and voiding in vias,
     forms stable solns, with good shelf life, is compatible with ArF
     photolithog., is applicable using any coating method, is capable of
     via fill and planarization, has good etching rates, and can be used
     in any semiconductor device fabrication process. An absorbing
     composition is described herein that includes at least one inorg .-
     based compound, at least one absorbing compound, and at least one
     material modification agent. Methods of making an absorbing
     composition are also described that includes: (a) combining at least
     one inorg.-based compound, at least one absorbing compound, at least
     one material modification agent, an acid/H2O mixture, and one or more
     solvents to form a reaction mixture; and (b) allowing the reaction
     mixture to form the absorbing composition at room temperature
     Another method of making an absorbing composition includes: (a)
     combining at least one inorg.-based compound, at least one absorbing
     compound, at least one material modification agent, an acid/H2O
     mixture, and one or more solvents to form a reaction mixture; and (b)
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heating the reaction mixture to form the absorbing composition Yet another method of making an absorbing composition is described that includes: (a) combining at least one inorg.-based compound, at least one absorbing compound, at least one material modification agent, and

one or more solvents to form a reaction mixture, wherein the at least one material modification agent comprises at least one acid and H2O; and (b) heating the reaction mixture to form an absorbing material, a coating or a film. In other methods of making an absorbing composition described herein, those methods include: (a) combining at least one inorg.-based compound, at least one absorbing compound, at least one material modification agent, and one or more solvents to form a reaction mixture, wherein the at least one material modification agent comprises at least one acid and H2O; and (b) allowing the reaction mixture to form an absorbing material, a coating or a film.

IT 313482-99-4 639088-18-9

RL: TEM (Technical or engineered material use); USES (Uses) (antireflective coatings for via fill and photolithog.

applications and

methods of preparation for device fabrication)

RN 313482-99-4 HCAPLUS

CN 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester (CA INDEX NAME)

RN 639088-18-9 HCAPLUS

CN 9-Phenanthrenecarboxylic acid, (triethoxysily1)methyl ester (CA INDEX

NAME)

IC ICM C08G CC 74-13 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) Section cross-reference(s): 76 71-43-2D, Benzene, reactive derivs. 72-48-0, Alizarin 75-59-2, ΙT Tetramethylammonium hydroxide 78-10-4, TEOS 81-64-1, Quinizarin 84-60-6, Anthraflavic acid 633-00-1, Rosolic acid 723-62-6, 9-Anthracenecarboxylic acid 780-69-8, Phenyltriethoxysilane 1343-98-2D, Silicic acid, derivs. 1468-95-7, 9-Anthracenemethanol 7440-21-3D, Silicon, compds. 8064-60-6, C.I. Direct Yellow 59 9003-53-6D, Polystyrene, derivs. 9005-12-3, Methylphenylsiloxane 10581-12-1, Tetramethylammonium acetate 16722-51-3, Tosylate, uses 25322-68-3, Polyethylene oxide 29355-26-8, Phenylazophenol 37114-85-5, Cetvltrimethylammonium nitrate 37181-39-8, Triflate 38542-94-8, Ammonium triflate 51374-75-5, Cetyltrimethylammonium acetate 79876-59-8, 2-Hydroxy-4-(3-triethoxysilylpropoxy)-diphenyl ketone 313482-99-4 442905-54-6 442905-55-7 639088-18-9 846606-04-0 RL: TEM (Technical or engineered material use); USES (Uses) (antireflective coatings for via fill and photolithog. applications and methods of preparation for device fabrication) L22 ANSWER 2 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN AN 2004:708028 HCAPLUS Full-text DN 142:269244 TT Antireflective spin coating material for photolithography IN Baldwin, Teresa; Kennedy, Joseph Honeywell International Inc., USA PA SO Taiwan, 10 pp. CODEN: TWXXA5 DT Patent Chinese LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE -------------------TW 570964 20040111 TW 2001-90131682 PI В 20011220 <--PRAI US 2001-335264P P 20011115 <--Antireflective coating materials for UV photolithog, include at least one absorbing compds. and at least one pH tuning agent that are incorporated into the materials which can be spin coated. Suitable

absorbing compds. are those that absorb around wavelengths such as 365 nm, 248 nm, 193 nm and 157 nm that maybe used in photolithog.

Suitable pH tuning agents not only adjust the pH of the final coating composition, but also influence the chemical performance and characteristics, mech. performance and structural makeup of the final spin-on composition that is part of the layered material, electronic component or semiconductor component, such that the final spin-on composition is more compatible with the resist material that is coupled to it. More specifically, the pH tuning agent strong influences the polymeric characteristics, the structural makeup and the spatial orientation that results in increasing the surface properties of the anti-reflective coating for optimal resist performance. In other words, a pH tuning agent that merely adjusts the pH of the spin-on material without influencing the mech. properties and structural makeup of the spin-on composition or the coupled resist material is not contemplated herein. A method of making absorbing and pH tuned spin-on materials includes combining at least one organic absorbing compound and at least one pH tuning agent with at least one silane reactant during synthesis of the spin-on materials and compns.

IT 639088-18-9

RL: MOA (Modifier or additive use); USES (Uses)

(antireflective spin coating material for photolithog.)

RN 639088-18-9 HCAPLUS

CN 9-Phenanthrenecarboxylic acid, (triethoxysily1) methyl ester (CA INDEX

NAME)

- IC ICM C09D005-00
- ${\tt CC}-{\tt 74-6}$ (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

Section cross-reference(s): 42

IT 81-64-1, 1,4-Dihydroxyanthraquinone 84-60-6 603-45-2 723-62-6, 9-Anthracenecarboxylic acid 780-69-8, Phenyltriethoxysilane 1468-

95-7, 9-Hydroxymethylanthracene 1689-82-3 5025-12-7 79876-59-8 442905-54-6 442905-55-7 639988-18-9 846606-03-9 846606-04-0

- RL: MOA (Modifier or additive use); USES (Uses) (antireflective spin coating material for photolithog.)
- L22 ANSWER 3 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN
- AN 2004:2893 HCAPLUS Full-text
- DN 140:59780
- TI Preparation of silyl alkyl esters of anthracene- and phenanthrene carboxylic acids as anti-reflective layers for photolithographic applications
- IN Lehmann, Lutz Uwe; Lonsky, Ralph
- PA Honeywell Specialty Chemicals Seelze G.m.b.H., Germany
- SO PCT Int. Appl., 40 pp.
- CODEN: PIXXD2
- DT Patent
- LA English FAN.CNT 1

| | | | | | KIND DATE | | | APPLICATION NO. | | | | | | DATE | |
|-----------------------------|-----|-----|-----|-----|-----------|-------------|------|-----------------|------------------|------|------|-----|------|------|------|
| - | | | | | _ | | | | | | | | | | |
| PI WO 2004 20030620 < | | 53 | | A1 | | 2003 | 1231 | | WO 2 | 003- | EP65 | 34 | | | |
| W: | | AG, | AL, | AM, | ΑT, | AU, | AZ, | BA, | BB, | BG, | BR, | BY, | BZ, | CA, | CH, |
| CN, | co, | CR, | CU, | CZ, | DE, | DK, | DM, | DZ, | EC, | EE, | ES, | FI, | GB, | GD, | GE, |
| GH, | GM. | HR. | HU. | TD. | TT | IN, | TS. | JP. | KE. | KG. | KP. | KR. | K7. | LC. | T.K. |
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| OM, | · | · | · | · | | MD, | | • | · | | · | · | · | · | · |
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| DW. | | | | | | UZ, MZ, | | | | | | | 2 W. | λM | h 7 |
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| TR, | FI, | FR, | GB, | GR, | HU, | IE, | ΙT, | LU, | MC, | NL, | PT, | RO, | SE, | SI, | SK, |
| | BF, | ВJ, | CF, | CG, | CI, | CM, | GΑ, | GN, | GQ, | GW, | ML, | MR, | ΝE, | SN, | TD, |
| TG DE 10227807 A1 | | | | | | A1 20040122 | | | DE 2002-10227807 | | | | | | |
| 20020621 < AU 2003242741 | | | | | | 2004 | 0106 | | AU 2 | 003- | 2427 | 41 | | | |
| 20030620 < EP 1539771 | | | | | | 2005 | 0615 | | EP 2 | 003- | 7606 | 60 | | | |
| 20030620 < | A1 | | | | | LI 2 | 005- | ,000 | 0,5 | | | | | | |
| EP 1539 | 771 | | | В1 | | 2006 | 1227 | | | | | | | | |

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT. IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK CN 1662546 20050831 CN 2003-814421 Α 20030620 <--JP 2005535616 T 20051124 JP 2004-514820 20030620 <--AT 349454 Т 20070115 AT 2003-760669 20030620 <--US 20060052569 US 2005-518060 A 1 20060309 20050906 <--PRAI DE 2002-10227807 А 20020621 <--WO 2003-EP6534 20030620 <--W OS CASREACT 140:59780; MARPAT 140:59780

GΙ

AB The invention relates to the preparation of silyl alkyl esters I (R1, R2, R3 = same or different alkyl, aryl, heteroaryl; R4, R5 = same or different H, halo, alkyl, aryl, heteroaryl; n = 1-10; R6 = halo, alkyl, aryl, heteroaryl, OH, alkoxy, aryl ether, (un)substituted amino, carboxy, carboxy, carboxylic amido, sulfonic acid ester, sulfonyl, thio, thioether, nitro, etc.; m = 0-4; T, X, Y, Z = C, a benzo group, which is substituted m-fold with R6 or unsubstituted, is condensed on the bonds T-X, X-Y, or Y-Z to form a trinuclear aromatic ring system, etc.), in particular of anthracene and phenanthrene carboxylic acids, a process for their preparation, compns. and polysiloxane compns. which contain the silyl alkyl esters and which can be used in particular in the semiconductor industry for the preparation of anti-reflective layers for photolithog. applications.

IT 313482-99-4P 639038-13-9P 639088-19-0P

RL: SPN (Synthetic preparation); TEM (Technical or engineered material $\ensuremath{\mathsf{TEM}}$

use); PREP (Preparation); USES (Uses)

(preparation of silyl alkyl esters of anthracene and phenanthrene carboxylic

acids as anti-reflective layers for photolithog. applications)

RN 313482-99-4 HCAPLUS

CN 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester (CA INDEX NAME)

RN 639088-18-9 HCAPLUS
CN 9-Phenanthrenecarboxylic acid, (triethoxysilyl)methyl ester (CA INDEX

NAME)

RN 639088-19-0 HCAPLUS

CN 9-Phenanthrenecarboxylic acid, 3-(triethoxysilyl)propyl ester (CA INDEX

NAME)

RN 639088-20-3 HCAPLUS

CN 9-Anthracenecarboxylic acid, 3-(trimethoxysily1)propyl ester (CA INDEX

NAME)

IC ICM C07F007-18

CC 29-7 (Organometallic and Organometalloidal Compounds)

Section cross-reference(s): 37, 76 T 313482-99-4P 639088-18-9P 639088-19-0P

IT 313482-99-4P 639088-18-9P 639088-19-0P 639088-20-3P

RL: SPN (Synthetic preparation); TEM (Technical or engineered material

use); PREP (Preparation); USES (Uses)

(preparation of silyl alkyl esters of anthracene and phenanthrene carboxylic

acids as anti-reflective layers for photolithog. applications)
RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 4 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN AN 2003:570019 HCAPLUS Full-text

DN 140:243463

 ${\tt TI} \quad {\tt Anthracene-organosiloxane \; spin-on \; antireflective \; coating \; for \; KrF} \\ {\tt lithography}$

AU Kennedy, Joseph; Baldwin-Hendricks, Teri; Hebert, Mello; Suedmeyer, Arlene

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CS
     Honeywell, Sunnyvale, CA, 94089, USA
SO
     Proceedings of SPIE-The International Society for Optical Engineering
(
     2003), 5039(Pt. 1, Advances in Resist Technology and Processing
     XX), 144-151
     CODEN: PSISDG; ISSN: 0277-786X
PB
     SPIE-The International Society for Optical Engineering
DT
     Journal
    English
LA
     A sacrificial, spin-on 248 nm UV absorbing organosiloxane film has
AB
     been developed to enable via first trench last (VFTL) dual-damascene
     patterning. Amongst other design objectives one key material
     requirement was that the film be SiO based to facilitate trench etch.
     Because the starting organosiloxane polymer is transparent a
     chromophores that absorbs at 248 nm had to be included. Anthracene
     was selected as it offered the largest amount of absorption per mol.
     Unfortunately, com. available anthracene moieties with different
     functional groups when added directly to the organosiloxane polymer
     solution resulted in films of very marginal quality. The primary
     issue was the poor solubility and thermal stability of the anthracene
     compound within the organosiloxane matrix. To address this
     fundamental problem the chromophore was stabilized by chemical
     attaching it to Et orthosilicate. The resulting mol., 9-anthracene
     carboxymethyltriethoxysilane (TESAC) was developed. By combining the
     appropriate amts. of TESAC, TEOS and other organo-TEOS monomers with
     the appropriate solvents and a catalyst a stable, 248 nm light
     absorbing anthracene-organosiloxane polymer was developed. This work
     has led to the development of Duo248 organosiloxane based bottom
     antireflective coatings. Topics such as the development of TESAC,
     lithog., plasma etch and selective removal will be discussed.
     650634-87-0
TT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (Duo 248; lithog, properties of anthracene-organosiloxane bottom
        antireflective coating for 248 nm lithog.)
RN
     650634-87-0 HCAPLUS
CN
     9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester, polymer
with
     silicic acid (H4SiO4) tetraethyl ester and silicic acid (H4SiO4)
triethyl
     methyl ester (CA INDEX NAME)
     CM
          1
     CRN 313482-99-4
     CMF C22 H26 O5 Si
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CRN 18395-48-7 CMF C7 H18 O4 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

IT 313482-99-4

RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent) (monomer; design of spin-on 248 nm UV absorbing anthraceneorganosiloxane film for bottom antireflective coatings)

RN 313482-99-4 HCAPLUS CN 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester (CA INDEX NAME)

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) Section cross-reference(s): 76

ΙT 650634-87-0

> RL: TEM (Technical or engineered material use); USES (Uses) (Duo 248; lithog, properties of anthracene-organosiloxane bottom antireflective coating for 248 nm lithog.)

313482-99-4 ΙT

RL: PRP (Properties); RCT (Reactant); RACT (Reactant or reagent) (monomer; design of spin-on 248 nm UV absorbing anthraceneorganosiloxane film for bottom antireflective coatings)

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 6 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 5 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:417795 HCAPLUS Full-text

DN 139:8200

TΙ Spin-on-glass antireflective coatings for photolithography

Baldwin-Hendricks, Teresa; Kennedy, Joe; Richey, Mary IN

PA Honeywell International Inc., USA

SO PCT Int. Appl., 63 pp., Added priority US2001-PV335259 w/ date 2001-11-15

to prevent further basic probl

CODEN: PIXXD2 Patent

DT T.A

| FAN.CNT 1 | | | | | | | | | | |
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| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE | | | | | |
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| PT | WO 2003044077 | A 1 | 20030530 | WO 2001-US43831 | | | | | | |

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W: JP, KP, KR, US

RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC,

NL,

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PT, SE, TR
     EP 1478681
                         A1 20041124 EP 2001-995897
20011116 <--
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
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             IE, FI, CY, TR
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                               20060821
                                          TW 2002-91133510
                        В
20021115 <--
    KR 818678
                        В1
                              20080401 KR 2004-707541
20040517 <--
PRAI US 2001-335259P
                         P
                               20011115 <--
    WO 2001-US43831
                         W
                               20011116 <--
     A siloxane polymer family comprising siloxane polymer made from (a) a
AB
     strongly UV absorbing phenylalkoxysilane, (b) ≥1 silane having good
     leaving groups, and (c) ≥1 silane having good leaving groups that is
     different than (b), where the siloxane polymer exhibits a nonlinear
     relationship, that is concave/convex or is located in the region
     enclosed by a concave/convex relation, between the ratio of
     (a):(b):(c) and the siloxane polymer's extinction coefficient k
     value. These siloxane polymers are preferably used as spin-on glass
     compns. for films in the microelectronics applications.
IΤ
     442905-62-6P, 9-Anthracenylcarboxypropyltriethoxysilane-
     methyltriethoxysilane-tetraethoxysilane copolymer
     RL: IMF (Industrial manufacture); PRP (Properties); TEM (Technical or
     engineered material use); PREP (Preparation); USES (Uses)
        (film; spin-on-glass antireflective coatings for mask used in
        microelectronics applications)
     442905-62-6 HCAPLUS
RN
CN
     9-Anthracenecarboxylic acid, 3-(triethoxysilyl)propyl ester, polymer
with
     silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane
(9CI)
     (CA INDEX NAME)
```

CM 1

CRN 442905-57-9

CMF C24 H30 05 Si

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

IT 313482-99-4P 442905-58-0P 442905-59-1P

RL: IMF (Industrial manufacture); PREP (Preparation)
 (spin-on-glass antireflective coatings for mask used in microelectronics applications)

RN 313482-99-4 HCAPLUS

CN 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester (CA INDEX NAME)

RN 442905-58-0 HCAPLUS

CN 9-Anthracenecarboxylic acid, 2-(triethoxysilyl)ethyl ester (CA INDEX NAME)

RN 442905-59-1 HCAPLUS

CN 9-Anthracenecarboxylic acid, (trimethoxysily1)methyl ester (CA INDEX NAME)

IT 442905-57-9P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation);

RACT

(Reactant or reagent)
(spin-on-glass antireflective coatings for mask used in

microelectronics applications)

RN 442905-57-9 HCAPLUS

CN 9-Anthracenecarboxylic acid, 3-(triethoxysilyl)propyl ester (CA INDEX

NAME)

IC ICM C08G077-02

CC 42-5 (Coatings, Inks, and Related Products)

Section cross-reference(s): 74, 76

IT 162746-16-9P, Phenyltriethoxysilane-methyltriethoxysilanetetraethoxysilane copolymer 442905-62-6P, 9-

Anthracenylcarboxypropyltriethoxysilane-methyltriethoxysilane-

tetraethoxysilane copolymer

RL: IMF (Industrial manufacture); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(film; spin-on-glass antireflective coatings for mask used in microelectronics applications)

II 313482-99-4P 442905-58-0P 442905-59-1P

RL: IMF (Industrial manufacture); PREP (Preparation)

(spin-on-glass antireflective coatings for mask used in

microelectronics applications)

IT 442905-57-9P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation);

RACT

(Reactant or reagent)

(spin-on-glass antireflective coatings for mask used in

microelectronics applications)

RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L22 ANSWER 6 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 2003:35308 HCAPLUS Full-text

DN 138:98200

TI Spin-on-glass anti-reflective coating for photolithography

IN Kennedy, Joseph; Baldwin, Teresa; Hacker, Nigel P.; Spear, Richard

PA Honeywell International Inc., USA

U.S., 13 pp., Cont.-in-part of U.S. 6,268,457. CODEN: USXXAM SO

DT Patent

LA English

| FAN.CNT 5 | | | | | | |
|-----------------------------|---------------|------------|-------------------------|-------------------|--|--|
| PATENT NO. | KIND | DATE | APPLICATION NO. | . DATE | | |
| | | | | | | |
| PI US 6506497 20000126 < | В1 | 20030114 | US 2000-491166 | | | |
| US 6268457 | В1 | 20010731 | US 1999-330248 | | | |
| 19990610 < CA 2374944 | A1 | 20001221 | CA 2000-2374944 | | | |
| 20000608 < | | | 011 2000 2071311 | | | |
| WO 2000077575 | A1 | 20001221 | WO 2000-US15772 | | | |
| 20000608 < | | | | | | |
| | , AU, AZ | , BA, BB, | BG, BR, BY, CA, CH, CN | , CU, CZ, | | |
| DE, DK. EE. ES | . FI. GB | . GE. GH. | GM, HU, ID, IL, IN, IS | JP. KE. | | |
| KG, | ,, | ,,, | ,,,,, | , , , , , , , , , | | |
| KP, KR, KZ | , LC, LK | , LR, LS, | LT, LU, LV, MD, MG, MK | , MN, MW, | | |
| MX, | | | | | | |
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| TT, | | | | | | |
| UA, UG, UZ | | | SL, SZ, TZ, UG, ZW, AT, | BF CH | | |
| CY, | , 113, FIN | , 142, 30, | 31, 32, 12, 09, 2W, AI | , DE, CII, | | |
| | , FI, FR | , GB, GR, | IE, IT, LU, MC, NL, PT | SE, BF, | | |
| BJ, | ,, | ,, | ,,,,, | ,, | | |
| CF, CG, CI | , CM, GA | , GN, GW, | ML, MR, NE, SN, TD, TG | | | |
| EP 1190277 | A1 | 20020327 | EP 2000-941275 | | | |
| 20000608 < | | | | | | |
| | , DE, DK | , ES, FR, | GB, GR, IT, LI, LU, NL | , SE, MC, | | |
| PT, | 7.17 ET | DO. | | | | |
| IE, SI, LT JP 2003502449 | , LV, FI Т | | JP 2001-503571 | | | |
| 20000608 < | 1 | 20030121 | OF 2001-303371 | | | |
| | A1 | 20020718 | US 2001-1143 | | | |
| 20011115 < | | | | | | |
| US 6824879 | B2 | 20041130 | | | | |
| US 20020128388 | A1 | 20020912 | US 2002-76846 | | | |
| 20020214 < | | | | | | |
| US 6956097 | В2 | 20051018 | | | | |
| US 20030120018 | A1 | 20030626 | US 2002-300357 | | | |
| 20021119 < | | | | | | |
| US 6969753 | B2 | 20051129 | HG 2005 130544 | | | |
| US 20050245717 | A1 | 20051103 | US 2005-178544 | | | |

| 2005071 | 1 < | | | |
|---------|--------------|----|----------|---------------|
| KR | 804873 | В1 | 20080220 | KR 2006-71552 |
| 2006073 | 1 < | | | |
| PRAI US | 1999-330248 | A2 | 19990610 | < |
| US | 2000-491166 | A | 20000126 | < |
| WO | 2000-US15772 | W | 20000608 | < |
| US | 2000-698883 | A2 | 20001027 | < |
| KR | 2001-715888 | A3 | 20011210 | < |
| US | 2002-76846 | A3 | 20020214 | < |
| US | 2002-300357 | A3 | 20021119 | < |

- AB Anti-reflective coating materials for deep UV photolithog. include one or more organic light-absorbing compds. incorporated into spin-on-glass materials. Suitable absorbing compds. are strongly absorbing over wavelength ranges around wavelengths such as 365 nm, 248 nm, and 193 nm that may be used in photolithog. A method of making absorbing spin-on-glass materials includes combining one or more organic absorbing compds. with alkoxysilane or halosilane reactants during synthesis of the spin-on-glass materials.
- IT 313482-99-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT

(Reactant or reagent)

(in preparation of spin-on-glass anti-reflective coatings for photolithog.)

RN 313482-99-4 HCAPLUS

CN 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester (CA INDEX NAME)

IT 442905-69-3P 442905-76-2P 442905-77-3P

RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(spin-on-glass anti-reflective coatings for photolithog.)

RN 442905-69-3 HCAPLUS CN 9-Anthracenecarboxyl:

CN 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane

(9CI)

(CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

RN 442905-76-2 HCAPLUS

 $\ensuremath{\mathsf{CN}}$ 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester, polymer with

trichloromethylsilane and trichlorosilane (9CI) (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM :

CRN 10025-78-2 CMF Cl3 H Si

CM 3

CRN 75-79-6 CMF C H3 Cl3 Si

RN 442905-77-3 HCAPLUS

CN 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester, polymer with

triethoxysilane (9CI) (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM 2

CRN 998-30-1 CMF C6 H16 O3 Si

OEt EtO-SiH-OEt

IC ICM B32B009-04

INCL 428447000; 528031000; 528034000; 528028000; 528026000; 528029000; 523137000; 106287110; 106287130; 106287140

 ${\tt CC} - 74-5$ (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

Section cross-reference(s): 38, 42 TT 313482-99-4P RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (in preparation of spin-on-glass anti-reflective coatings for photolithog.) IΤ 72-48-0DP, Alizarin, reaction products with silane derivs. 1468-95-7DP, 9-Anthracene methanol, reaction products with silane derivs. 88029-70-3DP, Methyltriethoxysilane-tetraethoxysilane copolymer, reaction products with 9-anthracene methanol 113923-94-7P, Phenvltriethoxysilanetetraethoxysilane copolymer 162746-16-9P, Methyltriethoxysilanephenyltriethoxysilane-tetraethoxysilane copolymer 442905-63-7P 442905-69-3P 442905-76-2P 442905-77-3P 442905-78-4P 484638-70-2DP, reaction products with 9-anthracene methanol and alizarine 484638-71-3DP, reaction products with 9-anthracene methanol and alizarin RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (spin-on-glass anti-reflective coatings for photolithog.) RE.CNT 55 THERE ARE 55 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT L22 ANSWER 7 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN AN 2002:540266 HCAPLUS Full-text 137:110586 DN TΤ Compositions, application, and spin-on-glass antireflective coatings for

photolithography

IN Baldwin, Teresa; Kennedy, Joseph; Hacker, Nigel; Spear, Richard

PA Honeywell International Inc., USA

SO U.S. Pat. Appl. Publ., 25 pp., Cont.-in-part of U.S. 6,268,457. CODEN: USXXCO

Patent DT

LA English

FAN.CNT 5 PATENT NO. KIND DATE APPLICATION NO. PI US 20020095018 A1 20020718 US 2001-1143 20011115 <--B2 20041130 US 6824879 US 6268457 В1 20010731 US 1999-330248

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19990610 <--
    US 6506497
                   B1
                              20030114 US 2000-491166
20000126 <--
                         R1
                                           US 2000-698883
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20001027 <--
    US 20020055000
                         A1
                               20020509
                                           US 2001-12649
20011105 <--
                         B2
    US 7012125
                               20060314
    WO 2003044079
                         A 1
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20021031 <--
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             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE,
GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
LR.
             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM,
PH.
             PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT,
TZ.
             UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ,
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                            20041124 EP 2002-773961
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    CN 1615333
                         A
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20021031 <--
    WO 2003044078 A1 20030530 WO 2002-US36327
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CN,
             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE,
GH.
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
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LR.
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PH.
             PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR,
TT.
             TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
BY.
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES,
             FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ,
CF,
             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
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                                20030610
                                           AU 2002-359387
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20021112 <--
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PT.
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK
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20021112 <--
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                         Α
                               20050511 CN 2002-827150
20021112 <--
PRAI US 1999-330248
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                               19990610 <--
     US 2000-491166
                          A2
                               20000126 <--
     US 2000-698883
                          A2
                               20001027 <--
     US 2001-1143
                          Α
                               20011115 <--
     WO 2001-US45306
                          А
                                20011115 <--
     WO 2002-US35101
                                20021031 <--
                          W
     WO 2002-US36327
                          W
                                20021112 <--
     Antireflective coating materials for UV photolithog. include ≥1
AB
     organic light-absorbing compound incorporated into spin-on-glass
     materials. Suitable absorbing compds. are strongly absorbing <375 nm
     that may be used in photolithog., where ≥1 of the silicon-based
     compound or the organic absorbing compound comprises an alkyl group,
     an alkoxy group, a ketone group, or an azo group. An additive of ≥1
     (reactive) organic absorbing compound is mixed with alkoxysilane or
     halosilane reactants during synthesis of the spin-on-glass materials.
     442905-57-9P 442905-58-0P 442905-59-1P
ΙT
     RL: IMF (Industrial manufacture); RCT (Reactant); TEM (Technical or
     engineered material use); PREP (Preparation); RACT (Reactant or
reagent);
     USES (Uses)
```

(light absorbent; spin-on-glass anti-reflective coatings for

photolithog.)

442905-57-9 HCAPLUS

RN

CN 9-Anthracenecarboxylic acid, 3-(triethoxysily1)propyl ester (CA INDEX

NAME)

RN 442905-58-0 HCAPLUS

CN 9-Anthracenecarboxylic acid, 2-(triethoxysilyl)ethyl ester (CA INDEX NAME)

RN 442905-59-1 HCAPLUS

CN 9-Anthracenecarboxylic acid, (trimethoxysily1)methyl ester (CA INDEX NAME)

IT 313482-99-4P, 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester

RL: IMF (Industrial manufacture); RCT (Reactant); TEM (Technical or engineered material use); PREP (Preparation); RACT (Reactant or reagent);

USES (Uses)

(spin-on-glass anti-reflective coatings for photolithog.) 313482-99-4 HCAPLUS

RN 313482-99-4 HCAPLUS

CN 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester (CA INDEX NAME)

IT 442905-61-5P 442905-62-6P 442905-65-9P 442905-66-0P 442905-67-1P 442905-68-2P

442905-69-3P 442905-70-6P 442905-71-7P

442905-73-9P 442905-75-1P 442905-76-2P

442965-77-3P

RL: IMF (Industrial manufacture); TEM (Technical or engineered material $\$

use); PREP (Preparation); USES (Uses)

(spin-on-glass coating; spin-on-glass anti-reflective coatings for photolithog.)

RN 442905-61-5 HCAPLUS

CN 9-Anthracenecarboxylic acid, 2-(trimethoxysily1)ethyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 7

CRN 442905-60-4

CMF C20 H22 O5 Si

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

RN 442905-62-6 HCAPLUS
CN 9-Anthracenecarboxylic acid, 3-(triethoxysilyl)propyl ester, polymer with silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI)

(CA INDEX NAME)

CRN 442905-57-9 CMF C24 H30 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, 5-(triethoxysily1)pentyl ester, polymer with }$

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-64-8 CMF C26 H34 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

RN 442905-66-0 HCAPLUS

 ${\tt CN}-{\tt 9-Anthracenecarboxylic}$ acid, (triethoxysily1)methy1 ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

CRN 681-84-5 CMF C4 H12 O4 Si

MeO-Si-OMe

RN 442905-67-1 HCAPLUS

CN 9-Anthracenecarboxylic acid, 2-(triethoxysily1)ethyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-58-0 CMF C23 H28 O5 Si

CM

CRN 1185-55-3 CMF C4 H12 O3 Si

CRN 681-84-5 CMF C4 H12 O4 Si

RN 442905-68-2 HCAPLUS CN

9-Anthracenecarboxylic acid, (trimethoxysilyl)methyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI)

(CA INDEX NAME)

CM 1

CRN 442905-59-1 CMF C19 H20 O5 Si

CRN 1185-55-3 CMF C4 H12 O3 Si

MeO-Si-Me

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

MeO—Si—OMe

RN 442905-69-3 HCAPLUS

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester, polymer with }$

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

RN 442905-70-6 HCAPLUS CN 9-Anthracenecarboxylic acid, 2-(triethoxysilyl)ethyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CRN 442905-58-0 CMF C23 H28 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, 3-(triethoxysily1)propyl ester, polymer with }$

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-57-9 CMF C24 H30 O5 Si

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

RN 442905-73-9 HCAPLUS

 \mbox{CN} 9-Anthracenecarboxylic acid, (tripropoxysily1)methyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-72-8 CMF C25 H32 O5 Si

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

CRN 681-84-5 CMF C4 H12 O4 Si

MeO-Si-OMe

RN 442905-75-1 HCAPLUS

CN 9-Anthracenecarboxylic acid, 2-(tributoxysily1)ethyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM

CRN 442905-74-0

CMF C29 H40 O5 Si

CM

CRN 1185-55-3

CMF C4 H12 03 Si

CRN 681-84-5 CMF C4 H12 O4 Si

RN 442905-76-2 HCAPLUS

 $\ensuremath{\mathsf{CN}}$ 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester, polymer with

trichloromethylsilane and trichlorosilane (9CI) (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

RN 442905-77-3 HCAPLUS

 $\ensuremath{\text{CN}}$ 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester, polymer with

triethoxysilane (9CI) (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

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CRN 998-30-1
    CMF C6 H16 O3 Si
     OEt
Eto-SiH-OEt
TC
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INCL 528039000
     42-5 (Coatings, Inks, and Related Products)
     Section cross-reference(s): 74
TΤ
    442905-57-9P 442905-58-0P 442905-59-1P
    RL: IMF (Industrial manufacture); RCT (Reactant); TEM (Technical or
     engineered material use); PREP (Preparation); RACT (Reactant or
reagent):
    USES (Uses)
        (light absorbent; spin-on-glass anti-reflective coatings for
       photolithog.)
ΙT
     313482-99-4P, 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl
     ester
     RL: IMF (Industrial manufacture); RCT (Reactant); TEM (Technical or
     engineered material use); PREP (Preparation); RACT (Reactant or
reagent);
    USES (Uses)
        (spin-on-glass anti-reflective coatings for photolithog.)
     88029-70-3P, Methyltriethoxysilane-tetraethoxysilane copolymer
TΤ
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     442905-70-6P 442905-71-7P 442905-73-9P
     442905-75-1P 442905-76-2P 442905-77-3P
     442905-78-4P
                   442905-79-5P
     RL: IMF (Industrial manufacture); TEM (Technical or engineered
material
     use); PREP (Preparation); USES (Uses)
        (spin-on-glass coating; spin-on-glass anti-reflective coatings for
       photolithog.)
RE.CNT 20
             THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
1.22 ANSWER 8 OF 8 HCAPLUS COPYRIGHT 2008 ACS on STN
     2000:900926 HCAPLUS Full-text
AN
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DN
    134:63904
TΙ
    Spin-on-glass anti-reflective coatings for photolithography
    Spear, Richard; Hacker, Nigel P.; Baldwin, Teresa; Kennedy, Joseph
IN
PA
    Alliedsignal Inc., USA
SO
    PCT Int. Appl., 38 pp.
    CODEN: PIXXD2
DT
    Patent
    English
LA
FAN.CNT 5
    PATENT NO.
                      KIND DATE
                                     APPLICATION NO.
                                                         DATE
    WO 2000077575
                       A1 20001221
PΤ
                                       WO 2000-US15772
20000608 <--
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DE.
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KG.
            KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW,
MX.
           NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR,
TT.
            UA, UG, UZ, VN, YU, ZW
       RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH,
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            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF,
BJ,
            CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                       B1 20010731 US 1999-330248
    US 6268457
19990610 <--
    US 6506497
                       B1 20030114 US 2000-491166
20000126 <--
    CA 2374944
                 A1 20001221 CA 2000-2374944
20000608 <--
    EP 1190277
                   A1 20020327 EP 2000-941275
20000608 <--
       R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
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                       T 20030121 JP 2001-503571
    JP 2003502449
20000608 <--
    KR 804873
                   B1 20080220 KR 2006-715525
20060731 <--
PRAI US 1999-330248 A
                            19990610 <--
    US 2000-491166
                            20000126 <--
                      A
                      W
    WO 2000-US15772
                             20000608 <--
    KR 2001-715888 A3
                            20011210 <--
```

- AB Anti-reflective coating materials for deep UV photolithog. include one or more organic light-absorbing compds. incorporated into spin-on-glass materials. Suitable absorbing compds. are strongly absorbing over wavelength ranges around wavelengths such as 365 nm, 248 nm, and 193 nm that may be used in photolithog. A method of making absorbing spin-on-glass materials includes combining one or more organic absorbing compds. with alkoxysilane or halosilane reactants during synthesis of the spin-on-glass materials.
- IT 313482-99-4

 RL: TEM (Technical or engineered material use); USES (Uses)

 (spin-on-qlass anti-reflective coatings for photolithog.)

RN 313482-99-4 HCAPLUS CN 9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester (CA INDEX NAME)

- IC ICM G03F007-09
 ICS H01L021-027; C08G077-14; C08G077-50; C07F007-18; C08K005-00; C09F183-04
- CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes) Section cross-reference(s): 42
- IT 72-48-0, Alizarin 81-64-1, Quinizarin 633-00-1, Rosolic acid 780-69-8, Phenyltriethoxysilane 1468-95-7, 9-Anthracenemethanol 79876-59-8 313482-99-4

RL: TEM (Technical or engineered material use); USES (Uses)
(spin-on-glass anti-reflective coatings for photolithog.)

E.CNT 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> D L23 1-12 BIB ABS HITSTR HITIND

L23 ANSWER 1 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 2005:490452 HCAPLUS Full-text

DN 143:51852

TI Selective removal chemistries for sacrificial layers, production and

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PA
     Honeywell International Inc., USA
     PCT Int. Appl., 35 pp.
SO
     CODEN: PIXXD2
DT
     Pat.ent.
LA
     English
FAN.CNT 1
     PATENT NO.
                         KIND
                                DATE
                                           APPLICATION NO.
                                                                   DATE
PT
     WO 2005053004
                         A1
                                20050609
                                            WO 2004-US38301
20041117 <--
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GD,
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LC,
             LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA,
NI,
             NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL,
SY,
             TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
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         RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW,
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DK,
             EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LU, MC, NL, PL, PT,
RO,
             SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
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             NE, SN, TD, TG
PRAI US 2003-523624P
                                20031119 <--
     Etching solns. described herein include (a) ≥1 fluorine-based
AB
     constituent, (b) \geq 1 acid constituent, such as a strong acid in some
     cases, and (c) a suitable solvent constituent, wherein the etching
     solution selectively etches at least one sacrificial layer from a
     surface or layered component. Methods for producing an etching
     solution include blending ≥1 fluorine-based constituent, acid
     constituent and compatible solvent constituent to form a solution,
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wherein the etching solution selectively etches at least one sacrificial layer from a surface or layered component. More particularly, the method of forming an etching solution includes blending hydrogen fluoride, hydrochloric acid as the acid and

propylene carbonate as the solvent.

uses IN

Starzynski, John S.

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IT 650634-87-0, DU0248
```

RL: REM (Removal or disposal); PROC (Process)
(fluoride compound-containing etching solns. for selective removal

of sacrificial layers)

RN 650634-87-0 HCAPLUS

 $\ensuremath{\mathsf{CN}}$ 9-Anthracenecarboxylic acid, (triethoxysily1)methyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester and silicic acid (H4SiO4) triethyl

methyl ester (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM 2

CRN 18395-48-7 CMF C7 H18 O4 Si

CM 3

CRN 78-10-4

Eto-Si-OEt

IC ICM H01L021-00

ICS H01L021-302

CC 76-3 (Electric Phenomena)

Section cross-reference(s): 74

RL: REM (Removal or disposal); PROC (Process)

(fluoride compound-containing etching solns. for selective removal

of sacrificial lavers)

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L23 ANSWER 2 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 2004:430850 HCAPLUS Full-text

DN 141:15725

 $\ensuremath{\mathsf{TI}}$ Antireflective silses quioxane coatings for photolithography and methods of

preparation thereof

- IN Baldwin, Teresa; Kennedy, Joseph; Iwamoto, Nancy; Nakano, Tadashi; Bedwell, William; Stuck, Jason; Suedemeyer, Arlene; Hebert, Mello; Li, Bo
- PA Honeywell International Inc., USA

SO PCT Int. Appl., 178 pp.

CODEN: PIXXD2

DT Patent

LA English

LA ENGLISH

| | PATENT NO. | | | | KINI |) - | DATE | | | APPLICATION NO. | | | | | | ATE |
|-----------------------|------------|--|----|--|------|--------|------|------|-----|-----------------|------|------|-----|----|----|------|
| - PI WO 2003111 | 2004 | | 25 | | A2 | | 2004 | 0527 | | WO 2 | 003- | US36 | 354 | | | |
| WO | 2004 | | | | A3 | 3 m | 2004 | | D.1 | DD | D.C. | D.D. | DV | DE | C. | OII. |

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,

CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE,

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GH.
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OM.
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TT,
             TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
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SK,
             TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
TD, TG
     WO 2003044600
                         A1
                                20030530 WO 2001-US45306
20011115 <--
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     AU 2002227106
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     CN 1606713
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                                           CN 2001-823932
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20011115 <--
     JP 2005512309
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                                20050428 JP 2003-546172
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WO 2003044078 A1 20030530 WO 2002-US36327
20021112 <--
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                         A9
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GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
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             LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM,
PH,
             PL, PT, RO, RU, SC, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR,
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             CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
     AU 2003295517
                         A1
                               20040603 AU 2003-295517
20031112 <--
     US 20050058929
                         A 1
                               20050317 US 2004-495688
20041117 <--
PRAI WO 2001-US45306
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    US 2002-444697P
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                               20021112 <--
    WO 2002-US36327
                         Α
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    US 2003-509199P
                         P
                               20031007 <--
    US 2001-1143
                         Α
                               20011115 <--
                               20031112 <--
    WO 2003-US36354
                         W
AB
     Antireflective coating materials for UV photolithog. in fabrication
     of semiconductor devices are based on silsesquioxanes having aromatic
     groups that absorb UV light around wavelengths such ad 365 nm, 248
     nm, 193 nm, and 157 nm. Optionally, these silsesquioxanes are
     manufactured in the presence of alkoxy silicates and have amine
     groups for control of pH. Optionally, UV absorbing compds. are added
     during the manufacture of of the silsesquioxanes. A typical
     silsesquioxane was manufactured by hydrolytic polymerization of TEOS
     2633.78, MeSi(OEt)3 1639.78, and 9-
     anthracenylcarbonyloxymethyltriethoxysilane 958.97 g.
```

IT 442905-62-6P, [3-(9-Anthracenylcarbonyloxy)propyl]triethoxysilanemethyltriethoxysilane-tetraethoxysilane copolymer 442905-69-3P, (9-Anthracenylcarbonyloxymethyl)triethoxysilane-

methyltriethoxysilane-

tetraethoxysilane copolymer 694509-45-0P, 3-Aminopropyltriethoxysilane-(9-

anthracenylcarbonyloxymethyl)triethoxysilane-

methyltriethoxysilane-tetraethoxysilane copolymer

RL: IMF (Industrial manufacture); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(antireflective silsesquioxane coatings for photolithog, in

fabrication of semiconductor devices)

RN 442905-62-6 HCAPLUS

 \mbox{CN} 9-Anthracenecarboxylic acid, 3-(triethoxysily1)propyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-57-9 CMF C24 H30 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

RN 442905-69-3 HCAPLUS

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester, polymer with$

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CRN 78-10-4 CMF C8 H20 O4 Si

RN 694509-45-0 HCAPLUS

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, (triethoxysilyl)methyl ester, polymer with$

silicic acid (H4SiO4) tetraethyl ester, triethoxymethylsilane and 3-(triethoxysilyl)-1-propanamine (9CI) (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CRN 919-30-2 CMF C9 H23 N O3 Si

CM 4

CRN 78-10-4 CMF C8 H20 O4 Si

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tetramethoxysilane copolymer 442905-67-1P, [2-(9-
     Anthracenvlcarbonvloxy)ethvlltriethoxysilane-methvltrimethoxysilane-
     tetramethoxysilane copolymer 442905-68-2P, (9-
     Anthracenylcarbonyloxymethyl)trimethoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 442905-70-6P, [2-(9-
     Anthracenylcarbonyloxy)ethyl]triethoxysilane-methyltriethoxysilane-
     tetraethoxysilane copolymer 442905-71-7P, [3-(9-
     Anthracenylcarbonyloxy)propyl]triethoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 442905-73-9P, (9-
     Anthracenylcarbonyloxymethyl)tripropoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 694509-47-2P, 3-
     Aminopropyltriethoxysilane-[2-(9-
anthracenvlcarbonvloxv)ethvlltriethoxvsil
     ane-methyltriethoxysilane-tetraethoxysilane copolymer 694509-48-3P
     , 3-Aminopropyltriethoxysilane-[3-(9-
anthracenvlcarbonvloxv)propvlltrietho
     xysilane-methyltriethoxysilane-tetraethoxysilane copolymer
     694509-50-7P, (9-Anthracenvlcarbonvloxymethyl)tributoxysilane-
     methyltrimethoxysilane-tetramethoxysilane copolymer
     RL: IMF (Industrial manufacture): TEM (Technical or engineered
material
     use); PREP (Preparation); USES (Uses)
        (antireflective silsesquioxane coatings for photolithog. in
fabrication
        of semiconductor devices)
     442905-65-9 HCAPLUS
     9-Anthracenecarboxylic acid, 5-(triethoxysilyl)pentyl ester, polymer
with
     silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane
(9CI)
     (CA INDEX NAME)
     CM
         1
     CRN 442905-64-8
     CMF C26 H34 O5 Si
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RN

CN

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CM 2

CRN 2031-67-6

CMF C7 H18 03 Si

OEt

OSI-Me

OET

CM 3

CRN 78-10-4

CMF C8 H20 04 Si

OET
```

(CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CRN 1185-55-3 CMF C4 H12 O3 Si

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

RN 442905-67-1 HCAPLUS
CN 9-Anthracenecarboxylic acid, 2-(triethoxysilyl)ethyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CRN 442905-58-0 CMF C23 H28 O5 Si

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, (trimethoxysily1)methyl ester, polymer with$

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-59-1 CMF C19 H20 O5 Si

CM 2

CRN 1185-55-3 CMF C4 H12 O3 Si

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

RN 442905-70-6 HCAPLUS

 $\mbox{CN} \quad \mbox{9-Anthracenecarboxylic acid, 2-(triethoxysilyl)ethyl ester, polymer with }$

silicic acid (H4SiO4) tetraethyl ester and triethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-58-0 CMF C23 H28 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CRN 78-10-4 CMF C8 H20 O4 Si

EtO—Si—OEt

RN 442905-71-7 HCAPLUS

CN 9-Anthracenecarboxylic acid, 3-(triethoxysily1)propyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 442905-57-9 CMF C24 H30 O5 Si

O OEt C-O-(CH2)3-Si-OEt

CM

CRN 1185-55-3

CMF C4 H12 03 Si

CRN 681-84-5 CMF C4 H12 O4 Si

RN 442905-73-9 HCAPLUS
CN 9-Anthracenecarboxylic acid, (tripropoxysilyl)methyl ester, polymer

with silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane

(9CI)

(CA INDEX NAME)

CM 1

CRN 442905-72-8 CMF C25 H32 O5 Si

CRN 1185-55-3 CMF C4 H12 O3 Si

MeO—Si—Me

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

MeO—Si—OMe

RN 694509-47-2 HCAPLUS

CN 9-Anthracenecarboxylic acid, 2-(triethoxysilyl)ethyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester, triethoxymethylsilane and 3-(triethoxysilyl)-1-propanamine (9CI) (CA INDEX NAME)

CM 1

CRN 442905-58-0 CMF C23 H28 O5 Si

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

CRN 919-30-2 CMF C9 H23 N O3 Si

CM 4

CRN 78-10-4 CMF C8 H20 O4 Si

RN 694509-48-3 HCAPLUS

CN 9-Anthracenecarboxylic acid, 3-(triethoxysily1)propyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester, triethoxymethylsilane and 3-(triethoxysilyl)-1-propanamine (9CI) (CA INDEX NAME)

CM 1

CRN 442905-57-9 CMF C24 H30 O5 Si

CM 2

CRN 2031-67-6 CMF C7 H18 O3 Si

CM 3

```
CRN 919-30-2
CMF C9 H23 N O3 Si
```

CRN 78-10-4 CMF C8 H20 O4 Si

RN 694509-50-7 HCAPLUS

CN 9-Anthracenecarboxylic acid, (tributoxysily1)methyl ester, polymer with

silicic acid (H4SiO4) tetramethyl ester and trimethoxymethylsilane (9CI) $\,$

(CA INDEX NAME)

CM 1

CRN 694509-49-4 CMF C28 H38 O5 Si

$$\bigcup_{i=0-CH_2-s}^{OBu-n}\bigcup_{i=0Bu-n}^{OBu-n}$$

CRN 1185-55-3 CMF C4 H12 O3 Si

CM 3

CRN 681-84-5 CMF C4 H12 O4 Si

- IC ICM C08G
- CC 76-3 (Electric Phenomena)
 - Section cross-reference(s): 74
- IT 88029-70-3P, Methyltriethoxysilane-tetraethoxysilane copolymer 442905-62-6P, [3-(9-Anthracenylcarbonyloxy)propyl]triethoxysilanemethyltriethoxysilane-tetraethoxysilane copolymer 442905-63-7P, 2-Hydroxy-4-[3-(triethoxysily])propoxy]diphenyl ketone-

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methyltriethoxysilane-tetraethoxysilane copolymer 442995-69-3P,
     (9-Anthracenylcarbonyloxymethyl)triethoxysilane-
methvltriethoxvsilane-
     tetraethoxysilane copolymer 694509-45-0P, 3-
     Aminopropyltriethoxysilane-(9-
anthracenylcarbonyloxymethyl)triethoxysilane-
     methyltriethoxysilane-tetraethoxysilane copolymer
     RL: IMF (Industrial manufacture); PRP (Properties); TEM (Technical or
     engineered material use); PREP (Preparation); USES (Uses)
        (antireflective silsesquioxane coatings for photolithog, in
fabrication
        of semiconductor devices)
     113923-94-7P, Phenyltriethoxysilane-tetraethoxysilane copolymer
ΙT
     175480-92-9P, 3-Aminopropyltriethoxysilane-methyltriethoxysilane-
     tetraethoxysilane copolymer 442905-65-9P, [5-(9-
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     tetraethoxysilane copolymer 442905-66-0P, (9-
     Anthracenylcarbonyloxymethyl)triethoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 442905-67-1P, [2-(9-
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     tetramethoxysilane copolymer 442905-68-2P, (9-
     Anthracenylcarbonyloxymethyl)trimethoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 442905-70-6P, [2-(9-
     Anthracenylcarbonyloxy)ethyl]triethoxysilane-methyltriethoxysilane-
     tetraethoxysilane copolymer 442905-71-7P, [3-(9-
     Anthracenylcarbonyloxy)propylltriethoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 442905-73-9P, (9-
     Anthracenylcarbonyloxymethyl)tripropoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 694509-46-1P, 3-
Aminopropyltriethoxysilane-
     2-hvdroxv-4-[3-(triethoxvsilvl)propoxvldiphenvl ketone-
     methyltriethoxysilane-tetraethoxysilane copolymer 694509-47-2P,
     3-Aminopropyltriethoxysilane-[2-(9-
anthracenylcarbonyloxy)ethyl]triethoxys
     ilane-methyltriethoxysilane-tetraethoxysilane copolymer
     694509-48-3P, 3-Aminopropyltriethoxysilane-[3-(9-
     anthracenylcarbonyloxy)propyl]triethoxysilane-methyltriethoxysilane-
     tetraethoxysilane copolymer 694509-50-7P, (9-
     Anthracenylcarbonyloxymethyl)tributoxysilane-methyltrimethoxysilane-
     tetramethoxysilane copolymer 694509-51-8P, 3-
Aminopropyltriethoxysilane-
     phenyltrimethoxysilane-tetraethoxysilane copolymer
                                                          694509-52-9P,
     3-Aminopropyltriethoxysilane-phenyltripropoxysilane-tetraethoxysilane
               694509-53-0P, 3-Aminopropyltriethoxysilane-
     phenyltributoxysilane-tetraethoxysilane copolymer 694509-54-1P,
     3-Aminopropyltriethoxysilane-4-ethoxy-4'-
(triethoxysilvlmethoxycarbonyl)az
```

```
obenzene-methyltriethoxysilane-tetraethoxysilane copolymer 694509-
56-3P.
     3-Aminopropyltriethoxysilane-4-ethoxy-4'-[2-
(triethoxysilyl)ethoxycarbonyl
     ]azobenzene-methyltriethoxysilane-tetraethoxysilane copolymer
     694509-58-5P, 3-Aminopropyltriethoxysilane-4-ethoxy-4'-[3-
     (triethoxysily1)propoxycarbonyl]azobenzene-methyltriethoxysilane-
     tetraethoxysilane copolymer 694509-59-6P, 4-Ethoxy-4'-[3-
     (triethoxysily1)propoxycarbonyl]azobenzene-methyltriethoxysilane-
     tetraethoxysilane copolymer 694509-60-9P, 3-
Aminopropyltriethoxysilane-
     methyltriethoxysilane-phenyltriethoxysilane-tetraethoxysilane
copolymer
    RL: IMF (Industrial manufacture); TEM (Technical or engineered
material
     use); PREP (Preparation); USES (Uses)
        (antireflective silsesquioxane coatings for photolithog. in
fabrication
       of semiconductor devices)
L23 ANSWER 3 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN
AN
     2004:80802 HCAPLUS Full-text
DN
    140:137660
    Composition and process for wet stripping removal of sacrificial
TI
    antireflective material
    Baum, Thomas H.; Bernhard, David; Minsek, David; Murphy, Melissa
IN
    Advanced Technology Materials, Inc., USA
PA
SO
    PCT Int. Appl., 39 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 1
    PATENT NO.
                 KIND DATE APPLICATION NO. DATE
PΙ
    WO 2004009730 A1 20040129 WO 2003-US22148
20030716 <--
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE,
GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ,
OM,
            PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,
TN.
```

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RW: GH. GM. KE. LS. MW. MZ. SD. SL. SZ. TZ. UG. ZM. ZW. AM. AZ.
BY.
             KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES.
             FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK,
TR,
             BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD,
TG
    US 20040016904
                         A 1
                                20040129
                                           US 2002-201340
20020723 <--
    US 6849200
                         B2
    AU 2003249279
                         A1
                                20040209
                                          AU 2003-249279
20030716 <--
    EP 1551936
                        A1 20050713 EP 2003-765600
20030716 <--
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
PT.
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
    CN 1671821
                         Α
                               20050921
                                           CN 2003-817397
20030716 <--
     JP 2005533896
                              20051110 JP 2004-523443
20030716 <--
PRAI US 2002-201340
                         A
                                20020723 <--
     WO 2003-US22148
                         W
                               20030716 <--
     The present invention relates to a wet stripping composition and
AB
     process useful for silicate stripping, e.g., wet stripping removal of
     sacrificial antireflective silicate material from a substrate or
     article having such material deposited thereon, particularly where
     the sacrificial antireflective silicate material is present with
     permanent silicate materials desired to be unaffected by the wet
     stripping composition The present invention in 1 aspect relates to a
     wet stripping composition comprising the following: a nitrogenous
     hydrofluoride; deionized H2O; organic solvent; and optionally a
     chelator and/or an amine/carboxylic acid buffer. The nitrogenous
     hydrofluoride may be of any suitable type, and comprises compds. such
     as NH4F and other amine hydrofluoride salts.
TТ
     650634-87-0
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC (Process);
```

TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW

(Uses)

USES

(etching of; composition and process for wet stripping removal of sacrificial antireflective material)

 $\rm RN~650634-87-0~HCAPLUS$ $\rm CN~9-Anthracenecarboxylic$ acid, (triethoxysilyl)methyl ester, polymer with

silicic acid (H4SiO4) tetraethyl ester and silicic acid (H4SiO4) triethyl $\,$

methyl ester (CA INDEX NAME)

CM 1

CRN 313482-99-4 CMF C22 H26 O5 Si

CM 2

CRN 18395-48-7 CMF C7 H18 O4 Si

CM 3

CRN 78-10-4 CMF C8 H20 O4 Si

```
ICS H01L021-302
    76-3 (Electric Phenomena)
CC
    Section cross-reference(s): 74
    78-10-4, Tetraethoxysilane 339984-98-4, Coral (barrier film)
IΤ
     477961-78-7, Trikon Orion 627863-28-9, Aurora 2.7 650594-25-5,
Duo 193
    650634-87-0 650634-88-1, Aurora 2.4
    RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); TEM (Technical or engineered material use); PROC (Process);
USES
     (Uses)
        (etching of; composition and process for wet stripping removal of
       sacrificial antireflective material)
RE.CNT 5
             THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L23 ANSWER 4 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN
AN
    2001:280495 HCAPLUS Full-text
DN
    134:318673
TI Positive-working photoresist composition for upper resist layer of
    composite two-layer resist
IN Uno, Seiji; Mizutani, Kazuvoshi
PA Fuji Photo Film Co., Ltd., Japan
SO Jpn. Kokai Tokkyo Koho, 58 pp.
    CODEN: JKXXAF
DT Patent
LA
    Japanese
FAN.CNT 1
    PATENT NO.
                 KIND DATE APPLICATION NO. DATE
PI JP 2001109150 A 20010420 JP 1999-284457
19991005 <--
PRAI JP 1999-284457
                              19991005 <--
AB
    The title composition contains acid-sensitive polysiloxane having
     repeating unit \{-Si[-L-X-Np(Z)](OA)m(Y-L-COOB)n]O3/2-\} (Np =
     naphthalene ring; A, B = H, acid-sensitive group; X = single bond; L
     = C1-10 alkylene, C3-12 cycloalkylene; X, Y = -OCO-; -COO-; -NHCO-;
     etc.; Z = halo, C1-10 alkyl; C3-12 cycloalkyl; C1-10 alkoxy, etc.;
     1, m, n = 0-3 integer) and a photoacid generator. The composition,
     which contains the polysiloxane, provides the improve storageability
     and generates little faulty resist.
IT 335262-51-6DP, 1-alkoxyethyl and pyranyl ethers
```

ICM C09K013-00

IC

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation);

(Reactant or reagent)

(polysiloxane in pos.-working photoresist composition)

RN 335262-51-6 HCAPLUS

 ${\tt CN} - 2 - {\tt Naphthalene} \mbox{carboxylic}$ acid, 6-hydroxy-, (triethoxysily1) methyl ester,

homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 335262-32-3 CMF C18 H24 O6 Si

IT 335262-34-5DP, 1-alkoxyethyl ether 335262-34-5P
335262-45-8DP, 1-alkoxyethyl ether 335262-49-2DP,

1-alkoxyethyl ether 335262-53-8DP, 1-alkoxyethyl ethers

335262-56-1DP, 1-alkoxyethyl ether 335277-03-7P 335277-06-0P

RL: SPN (Synthetic preparation); TEM (Technical or engineered material

use); PREP (Preparation); USES (Uses)

(polysiloxane in pos.-working photoresist composition)

RN 335262-34-5 HCAPLUS

 ${\tt CN} - 2{\tt -Naphthalenecarboxylic}$ acid, 6-hydroxy-, (triethoxysily1)methyl ester,

polymer with bicyclo[2.2.1]hept-2-ylsilanetriol (9CI) (CA INDEX NAME)

CM 1

CRN 335262-33-4 CMF C7 H14 O3 Si

CRN 335262-32-3 CMF C18 H24 O6 Si

RN 335262-34-5 HCAPLUS

 ${\tt CN} - 2-{\tt Naphthalenecarboxylic}$ acid, 6-hydroxy-, (triethoxysily1)methyl ester,

polymer with bicyclo[2.2.1]hept-2-ylsilanetriol (9CI) (CA INDEX NAME)

CM

CRN 335262-33-4 CMF C7 H14 O3 Si

CRN 335262-32-3 CMF C18 H24 O6 Si

RN 335262-45-8 HCAPLUS
CN 2-Naphthalenecarboxylic acid, 6-hydroxy-, (triethoxysilyl)methyl
ester,

polymer with cyclohexylsilanetriol (9CI) (CA INDEX NAME)

CM 1

CRN 335262-32-3 CMF C18 H24 O6 Si

CM 2

CRN 3553-74-0 CMF C6 H14 O3 Si

RN 335262-49-2 HCAPLUS

2-Naphthalenecarboxylic acid, 3,7-dihydroxy-, (triethoxysily1)methyl ester, polymer with cyclohexylsilanetriol (9CI) (CA INDEX NAME)

CM 1

CN

CRN 335262-48-1 CMF C18 H24 O7 Si

$$\begin{array}{c} \text{ODED} \\ \text{HO} \\ \text{OH} \end{array} \\ \begin{array}{c} \text{OET} \\ \text{OET} \\ \text{OH} \end{array}$$

CM 2

CRN 3553-74-0 CMF C6 H14 O3 Si

RN 335262-53-8 HCAPLUS CN 2-Naphthalenecarboxy

2-Naphthalenecarboxylic acid, 3,7-dihydroxy-, (triethoxysilyl)methyl ester, homopolymer (9CI) (CA INDEX NAME)

CRN 335262-48-1 CMF C18 H24 O7 Si

$$\begin{array}{c} \text{ODE} \\ \text{CO-CH2-} \\ \text{OH} \end{array}$$

RN 335262-56-1 HCAPLUS

CN 2-Naphthalenecarboxylic acid, 3-hydroxy-, 3-(triethoxysily1)propyl ester,

homopolymer (9CI) (CA INDEX NAME)

CM

1

CRN 335262-55-0 CMF C20 H28 O6 Si

RN 335277-03-7 HCAPLUS

CN 2,6-Naphthalenedicarboxylic acid, mono[(triethoxysily1)methyl] ester, homopolymer, 1-ethoxyethyl ester (9CI) (CA INDEX NAME)

CM 1

CRN 7518-70-9

CMF C4 H10 O2

CRN 335277-02-6

CMF (C19 H24 O7 Si)x

CCI PMS

CM 3

CRN 335277-01-5 CMF C19 H24 O7 Si

RN 335277-06-0 HCAPLUS

2,3-Naphthalenedicarboxylic acid, mono[(triethoxysily1)methyl] ester, homopolymer, 1-(2-methylpropoxy)ethyl ester (9CI) (CA INDEX NAME)

CM 1

CN

CRN 205813-86-1 CMF C6 H14 O2

i-Buo-CH-Mo

CM 2

CRN 335277-05-9 CMF (C19 H24 O7 Si)x CCI PMS

CM 3

CRN 335277-04-8 CMF C19 H24 O7 Si

ICM G03F007-039

335277-12-8P

material

IC

```
ICS C08L083-06; G03F007-075; G03F007-095; G03F007-26
    74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other
    Reprographic Processes)
    Section cross-reference(s): 76
ΤТ
    335262-51-6DP, 1-alkoxyethyl and pyranyl ethers
    RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation);
RACT
    (Reactant or reagent)
        (polysiloxane in pos.-working photoresist composition)
ΙT
    109-53-5DP, hydroxyaryl silsesquioxane ethers 110-87-2DP,
hydroxyaryl
    silsesquioxane ethers 335262-24-3DP, ethers 335262-24-3P
    335262-26-5P 335262-28-7P 335262-30-1P 335262-34-5DP,
    1-alkoxyethyl ether 335262-34-5P 335262-37-8DP, ethers
    335262-39-0DP, ethers 335262-41-4DP, ethers 335262-43-6DP, ethers
    335262-45-8DP, 1-alkoxyethyl ether 335262-49-2DP,
    1-alkoxyethyl ether 335262-53-8DP, 1-alkoxyethyl ethers
    335262-56-1DP, 1-alkoxyethyl ether 335262-59-4DP, 1-alkoxyethyl
```

ether 335262-61-8DP, 1-alkoxyethyl ether 335277-00-4P 335277-03-7P 335277-06-0P 335277-08-2P 335277-10-6P

RL: SPN (Synthetic preparation); TEM (Technical or engineered

(polysiloxane in pos.-working photoresist composition)

L23 ANSWER 5 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

use); PREP (Preparation); USES (Uses)

AN 1992:128093 HCAPLUS Full-text

DN 116:128093

OREF 116:21667a,21670a

 $\ensuremath{\mathsf{TI}}$ $\ensuremath{\mathsf{Effect}}$ of superfluous remote polar functionality on chiral recognition

AU Pirkle, William H.; Welch, Christopher J.

CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA

SO Journal of Chromatography (1992), 589(1-2), 45-51

CODEN: JOCRAM; ISSN: 0021-9673

DT Journal

LA English

AB During the liquid chromatog. separation of enantiomers on chiral stationary phases (CSFs), nonspecific adsorption processes diminish the observed enantioselectivity. The role of those polar groups in the analyte which are not specifically required for chiral recognition was investigated. A series of racemic analytes bearing various nonessential polar groups spatially removed from the sites of chiral recognition were synthesized and chromatog. evaluated. Four scenarios for the interaction of these remote polar groups with the CSP were considered and used to rationalize exptl. findings. Cases were observed where enantioselectivity decreased, increased or remained the same on incorporation of a remote polar group.

IT 99727-12-5D, reaction product with silica gel 130859-81-3D
, reaction product with silica gel

RL: PRP (Properties)

(chiral stationary phase, for liquid chromatog. separation of enantiomers) $% \left(\frac{1}{2}\right) =\frac{1}{2}\left(\frac{1}{2}\right) +\frac{1}{2}\left(\frac{1}{2}\right) +$

RN 99727-12-5 HCAPLUS

CN L-Valine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

RN 130859-81-3 HCAPLUS

CN D-Alanine, N-2-naphthalenyl-, l1-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

CC 22-13 (Physical Organic Chemistry) Section cross-reference(s): 34, 80

IT 90697-04-4D, reaction product with silica gel 95544-22-2D, reaction product with silica gel 99727-12-5D, reaction product with silica gel 130859-81-3D, reaction product with silica gel 139456-46-5D, reaction product with silica gel 139456-47-6D,

reaction

product with silica gel 139456-48-7D, reaction product with silica

gel

139456-49-8D, reaction product with silica gel RL: PRP (Properties)

(chiral stationary phase, for liquid chromatog. separation of enantiomers) $% \left(1\right) =\left(1\right) \left(1\right)$

L23 ANSWER 6 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 1991:680494 HCAPLUS Full-text

DN 115:280494

OREF 115:47687a,47690a

- TI A chiral stationary phase which affords unusually high levels of enantioselectivity
- AU Pirkle, William H.; Deming, Kris C.; Burke, John A., III
- CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA
- SO Chirality (1991), 3(3), 183-7 CODEN: CHRLEP; ISSN: 0899-0042

DT Journal

LA English

OS CASREACT 115:280494

GΙ

AB A chiral stationary phase (CSP) I derived from N-(1-naphthyl)leucine has been prepared This CSP is conceptually similar to the CSP derived from N-(2-naphthyl)alanine and was expected to sep. the enantiomers of the same clientele of analytes as does the latter. The magnitudes of the separation factors observed on the two CSPs may differ markedly for a given analyte, the new CSP often affording much greater enantioselectivity.

IT 130859-81-3

RL: RCT (Reactant); RACT (Reactant or reagent)
(chiral stationary phase derived from)

RN 130859-81-3 HCAPLUS

CN D-Alanine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA

UN D-Alanine, N-z-naphthalenyi-, ii-(triethoxysiiyi)undecyi ester (UINDEX NAME)

Absolute stereochemistry.

IT 137046-42-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation);

RACT

(Reactant or reagent)

(preparation and reaction of, with silica)

RN 137046-42-5 HCAPLUS

CN L-Leucine, N-1-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

IT 137046-42-50P, reaction product with silica

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of, as chiral stationary phase for separation of enantiomers)

RN 137046-42-5 HCAPLUS

CN L-Leucine, N-1-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA TNDEX

NAME)

Absolute stereochemistry.

CC 34-2 (Amino Acids, Peptides, and Proteins)

Section cross-reference(s): 25

IT 130859-81-3

RL: RCT (Reactant); RACT (Reactant or reagent)

```
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation);
RACT
     (Reactant or reagent)
        (preparation and reaction of, with silica)
ΙT
     137046-42-5DP, reaction product with silica
     RL: SPN (Synthetic preparation); PREP (Preparation)
        (preparation of, as chiral stationary phase for separation of
enantiomers)
L23 ANSWER 7 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN
AN
     1991:632788 HCAPLUS Full-text
DN
     115:232788
OREF 115:39705a,39708a
TT
    Fast liquid chromatography for the resolution of chiral compounds
AU
     Perrin, Scott R.
CS
     Regis Chem. Co., Morton Grove, IL, 60053, USA
SO
    Chirality (1991), 3(3), 188-95
     CODEN: CHRLEP: ISSN: 0899-0042
DT
    Journal
LA
    English
AB
    A Pirkle-concept chiral stationary phase (CSP) derived from N-(1-
     naphthyl) leucine was evaluated for developing methods to reduce anal.
     times and investigating techniques in the rapid screening of a
     variety of chiral compds., e.g. N-(3,5-dinitrobenzoyl)-DL-leucine
     amides, over a given chiral selector. The effects of reduced column
     lengths and elevated temps. were studied to shorten anal. times.
     137046-42-5D, silica-bound
TΤ
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (as chiral stationary phase for fast liquid chromatog, resolution
of chiral
        compds.)
RN
     137046-42-5 HCAPLUS
CN
     L-Leucine, N-1-naphthalenvl-, 11-(triethoxysilvl)undecvl ester (CA
INDEX
     NAME)
Absolute stereochemistry.
```

(chiral stationary phase derived from)

TT

137046-42-5P

CC 34-2 (Amino Acids, Peptides, and Proteins)
IT 137046-42-5D, silica-bound
RL: RCT (Reactant); RACT (Reactant or reagent)
(as chiral stationary phase for fast liquid chromatog. resolution of chiral

compds.)

L23 ANSWER 8 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN AN 1991:573805 HCAPLUS Full-text

AN 1991:573805 DN 115:173805

OREF 115:29443a,29446a

TI The influence of end-capping on the enantioselectivity of a chiral phase

AU Pirkle, W. H.; Readnour, R. S.

CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA

SO Chromatographia (1991), 31(3-4), 129-32

CODEN: CHRGB7; ISSN: 0009-5893

DT Journal

LA English

AB The effect of end-capping chiral stationary phases derived from N-(2-naphthyl)alanine undecyl ester has been examined using either trimethylchlorosilane (TMCS), hexamethyldisilazane (HMDS), or bis(trimethylsilyl)trifluoroacetamide (BSTFA) as end-capping reagents. The separation factor (α) and capacity factor (k') of the enantiomers of N-(3,5-dinitrobenzoyl)leucine octadecyl amide and N-(3,5-dinitrobenzoyl)leucine octadecyl amide and N-(3,5-dinitrobenzoyl)alanine Bu ester were evaluated on three columns all packed with material from the same batch of stationary phase. These columns were essentially identical before, but not after end-capping with the above reagents. TMCS and HMDS were found to be superior to BSTFA, which appears to cause a significant loss of bonded phase from the silica surface. It seems that residual silanols affect the retention either by interacting with the analyte or by interacting with strands of stationary phase. End-capping

usually increases enantioselectivity, sometimes by decreasing k' for the first enantiomer and increasing k' for the second enantiomer. The enhancement in enantioselectivity is greatest in relatively nonpolar mobile phases and occurs to a greater extent for phases having incomplete surface coverages.

ΙT 104336-53-0D, derivs. RL: ANST (Analytical study)

(as chiral stationary phase, in liquid chromatog. separation of enantiomers)

104336-53-0 HCAPLUS RN

L-Alanine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA CN INDEX

NAME)

Absolute stereochemistry.

CC 80-3 (Organic Analytical Chemistry) Section cross-reference(s): 34

ΙT 75-77-4D, Trimethylchlorosilane, reaction products with triethylamine and

naphthylalanine derivs. 121-44-8D, Triethylamine, reaction products with

trimethylchlorosilane and naphthylalanine derivs. Hexamethyldisilazane, reaction products with naphthylalanine derivs. 25561-30-2D, Bis(trimethylsilyl)trifluoroacetamide, reaction products

with naphthylalanine derivs. 104336-53-0D, derivs.

RL: ANST (Analytical study)

(as chiral stationary phase, in liquid chromatog. separation of enantiomers)

L23 ANSWER 9 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 1991:12758 HCAPLUS Full-text

DN 114:12758

OREF 114:2227a,2230a

TΤ Chromatographic approach to the measurement of the interstrand distance

for some chiral bonded phases

- AU Pirkle, William H.; Readnour, Robin S.
- CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA
- SO Analytical Chemistry (1991), 63(1), 16-20 CODEN: ANCHAM: ISSN: 0003-2700
- DT Journal
- LA English
- AB A series of homologous N, N'-bis (2, 4-dinitrophenyl) $-\alpha$, ω -

diaminoalkanes (di-DNP) was chromatographed at various temps. on π basic chiral stationary phases derived from N-(2-naphthyl)alanine to determine the enthalpy and entropy of adsorption. The number of CH2 groups in the di-DNP influences the ability of the terminal π -acidic groups to interact simultaneously with neighboring strands of stationary phase, a process termed "bridging". When the number of CH2 groups is optimal for bridging, the enthalpy of adsorption is most exothermic. The length of the dianalyte required for optimal bridging is related to the interstrand distance. Optimal bridging occurs for the di-DNP having 5 CH2 groups regardless of the extent of surface coverage of the silica for the surface coverage range investigated. This suggests that the strands are not randomly spaced on the silica, with interstrand distance being influenced only by surface coverage, but are instead clustered, the clusters having similar distributions of interstrand distances. Adsorption is more exothermic for phases of high surface coverages than for low. If interstrand spacing is independent of surface coverage but surface coverage affects the enthalpy of adsorption, then surface coverage must influence the cluster size, which then influences the average extent of solvation of a strand of bonded phase.

IT 104336-53-0D, reaction products with silica gel 130859-81-3D, reaction products with silica gel RL: PRP (Properties)

(surface, chromatog. chiral bonded phase, interstrand distance of) 104336-53-0 HCAPLUS

RN 104336-53-0 HCAPLUS CN L-Alanine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

RN 130859-81-3 HCAPLUS CN D-Alanine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX NAME)

Absolute stereochemistry.

CC 66-4 (Surface Chemistry and Colloids) Section cross-reference(s): 22, 69

IT 104335-53-0D, reaction products with silica gel 130859-81-3D, reaction products with silica gel RL: PRP (Properties)

(surface, chromatog. chiral bonded phase, interstrand distance of)

L23 ANSWER 10 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 1987:226783 HCAPLUS Full-text

DN 106:226783

OREF 106:36589a,36592a

TI Direct chromatographic separation of enantiomeric diol derivatives

AU Pirkle, W. H.; Mahler, George S.; Pochapsky, Thomas C.; Hyun, Myung Ho

CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA

SO Journal of Chromatography (1987), 388(2), 307-14

CODEN: JOCRAM; ISSN: 0021-9673

DT Journal

LA English

AB The enantiomers of a variety of aliphatic and alicyclic vicinal diols, derivatized as the bis(3,5-dinitrophenyl carbamates), were separated by liquid chromatog, on chiral stationary phases derived from either (R)-N-(2-naphthyl)alanine or an (R)-N-acylated α -aryl- α -aminoalkane. The technique has utility not only for anal detns. of enantiomeric purity and absolute configuration but also for preparative sepns. since the diols can be easily recovered from the separated derivs. Chiral recognition mechanisms are presented to account for the observed sepns.

IT 164336-53-0D, reaction products with silica gel RL: ANST (Analytical study)

(as stationary phases, in liquid chromatog. separation of

enantiomeric diols

as dinitrophenyl carbamate derivs.)

RN 104336-53-0 HCAPLUS

CN L-Alanine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

CC 80-4 (Organic Analytical Chemistry)

Section cross-reference(s): 23, 24

IT 95544-22-2D, reaction products with silica gel 104336-53-0D, reaction products with silica gel

RL: ANST (Analytical study)

(as stationary phases, in liquid chromatog. separation of enantiomeric diols

as dinitrophenyl carbamate derivs.)

L23 ANSWER 11 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 1987:137657 HCAPLUS Full-text

DN 106:137657

OREF 106:22447a,22448a

TI Useful and easily prepared chiral stationary phases for the direct chromatographic separation of the enantiomers of a variety of derivatized

amines, amino acids, alcohols, and related compounds

AU Pirkle, William H.; Pochapsky, Thomas C.; Mahler, George S.; Corey, Debbi

E.; Reno, Daniel S.; Alessi, Donna M.

CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA

SO Journal of Organic Chemistry (1986), 51(25), 4991-5000 CODEN: JOCEAH; ISSN: 0022-3263

DT Journal

LA English

OS CASREACT 106:137657

AB Chiral stationary phases (CSPs) derived from 10-undecenyl esters of N-(2-naphthyl)- α -amino acids were prepared from readily available enantiomerically pure α -amino acids. Such CSPs are useful for the

chromatog. separation of the enantiomers of a variety of functionalized chiral mols. including many amines, alcs., $\alpha-$ and $\beta-$ amino acids, and related compds., all as their 3,5-dinitrobenzamides, (3,5- dinitroanilino)ureas, or (3,5-dinitroanilino)carbamates. Separability factors exceeding 18 at room temperature were observed in some instances. A simple chiral recognition model is proposed to account for the remarkable regular order of elution of enantiomers from these CSPs.

IT 99727-12-5P 104336-53-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and immobilization on silica gel)

RN 99727-12-5 HCAPLUS

CN L-Valine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

RN 104336-53-0 HCAPLUS

CN L-Alanine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and immobilization on silica gel)

L23 ANSWER 12 OF 12 HCAPLUS COPYRIGHT 2008 ACS on STN

AN 1986:88951 HCAPLUS Full-text

DN 104:88951

OREF 104:14135a,14138a

TI A new, easily accessible reciprocal chiral stationary phase for the chromatographic separation of enantiomers

AU Pirkle, W. H.; Pochapsky, Thomas C.

CS Sch. Chem. Sci., Univ. Illinois, Urbana, IL, 61801, USA

SO Journal of the American Chemical Society (1986), 108(2), 352-4 CODEN: JACSAT; ISSN: 0002-7863

DT Journal LA English

LA English
OS CASREACT 104:88951

OS CASREACT 104:88951

AB A chiral stationary phase (CSP) consisting of (S)-(-)-ωsiloxyundecanyl N-(2-naphthyl)valinate covalently linked to
microparticulate silica is effective for the direct chromatog.
separation of the enantiomers of chiral amine and amino acid 3,5dinitrobenzamides, 3,5-dinitroanilinocarbamates derived from chiral
alcs. and thiols, and chiral amines as their 3,5-dinitroanilinourea
derivs. Twenty resolns. are presented. The CSP may be synthesized
from L-valine. Chromatog. efficiencies are comparable to typical
HPLC silica-bonded phases, and the CSP can be used in either a normal
or reverse-phase fashion.

IT 99727-12-5P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT

(Reactant or reagent)

(preparation and reaction of, with silica)

RN 99727-12-5 HCAPLUS

CN L-Valine, N-2-naphthalenyl-, 11-(triethoxysilyl)undecyl ester (CA INDEX

NAME)

Absolute stereochemistry.

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CC 34-2 (Amino Acids, Peptides, and Proteins)
Section cross-reference(s): 80

IT 9572-12-5P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation);
RACT
(Reactant or reagent)
(preparation and reaction of, with silica)
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